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(54) Title: PYRROLO-PYRAZINE DERIVATIVES USEFUL AS CB1-MODULATORS

$$\begin{array}{c|c}
R^2 & N & R^3 \\
\hline
R^1 & N & X
\end{array}$$
(I)

(57) Abstract: The present invention relates to compounds of formula (I); and pharmaceutically acceptable salts thereof, in which R¹ and R² independently represent phenyl, thienyl, pyridyl, C₁-10alkyl, C₁-10alkoxy or C₃-15cycloalkyl; R³ represents a C₁-15alkyl group, C₃-15cycloalkyl, a phenylC₁-4alkyl group, a heteroaryl group, a heteroarylC₁-4alkyl group, or a group R⁴(CH₂)n- in which R⁴ represents a saturated or partially unsaturated 5 to 8 membered heterocyclic group containing one or more heteroatoms selected from nitrogen, oxygen or sulphur and n is 0, 1, 2, 3 or 4; X and Y independently represent O or S; m and n independently represent 0 or 1; wherein each of R¹, R², R³ and R⁴ is optionally substituted by one, two or three groups represented by Z wherein Z represents a C₁-6alkyl group optionally substituted by one or more fluoro, a C₁-6alkoxy group optionally substituted by one or more fluoro, hydroxy, halo, trifluoromethylsulphonyl, benzyl, nitro, amino, mono or di C₁-4alkylamino, mono or di C₁-3alkylamido, C₁-3alkylsulphonyl, C₁-6alkoxycarbonyl, carboxy, cyano, carbamoyl, mono or di C₁-3alkyl carbamoyl, sulphamoyl or acetyl and processes for preparing such compounds, their use in the treatment of obesity, psychiatric and neurological disorders, to methods for their therapeutic use and to pharmaceutical compositions containing them.